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INFORMATION DISCLOSURE STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number	10571991
Filing Date	2006-03-15
First Named Inventor	Bernard Barlaam
Art Unit	1624
Examiner Name	Douglas M. Willis
Attorney Docket Number	09963.0008

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	1	5576322		1996-11-19	Takase et al.	

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	1	20040053972	A1	2004-03-18	Nara et al.	
	2	20040192664	A1	2004-09-30	Kunz et al.	
	3	20040198997	A1	2004-10-07	Scholz et al.	
	4	20050130995	A1	2005-06-16	Nishino et al.	
	5	20050148607	A1	2005-07-07	Suzuki et al.	

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	6	20060167026	A1	2006-07-27	Nawa et al.	
	7	20060188501	A1	2006-08-24	Homma et al.	
	8	20090312313	A1	2009-12-17	Shimizu et al.	

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	1	2001/19788	WO	A2	2001-03-22	COR Therapeutics, Inc.	(Submitted in 3 parts)	☐
	2	2001/21160	WO	A2	2001-03-29	Axxima Pharmaceuticals Aktiengesellschaft		☐
	3	2001/32155	WO	A2	2001-05-10	The University of Manchester		☐
	4	2001/64642	WO	A2	2001-09-07	COR Therapeutics, Inc.	(Submitted in 3 parts)	☐
	5	2002/05791	WO	A2	2002-01-24	Pharmacia & Upjohn S.P.A.		☐
	6	2002/17712	WO	A2	2002-03-07	FMC Corporation	(Submitted in 2 parts)	☐

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	7	2002/20020	WO	A1	2002-03-14	Pharmacia & Upjohn, S.P.A.		☐
	8	2002/30358	WO	A2	2002-04-18	Tularik, Inc.		☐
	9	2003/031406	WO	A2	2003-04-17	IRM LLC		☐
	10	2003/097086	WO	A2	2003-11-27	Technische Universitat Munchen	(Submitted in 2 parts)	☐
	11	2003/097615	WO	A1	2003-11-27	Scios, Inc.	(Submitted in 2 parts)	☐
	12	2003/099276	WO	A1	2003-12-04	Bristol-Myers Squibb Company	(Submitted in 7 parts)	☐
	13	2004/010929	WO	A2	2004-02-05	Scios, Inc.		☐
	14	2004/072038	WO	A1	2004-08-26	Vertex Pharmaceuticals, Inc.		☐
	15	2004/085385	WO	A2	2004-10-07	Schering Corporation		☐
	16	2004/096224	WO	A2	2004-11-11	Boehringer Ingelheim International GmbH		☐
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	18	2005/003325	WO	A2	2005-01-13	Dana Farber Cancer Institute		☐
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	20	2005/030140	WO	A2	2005-04-07	Ex-Elixis, Inc.	(Submitted in 4 parts)	☐
	21	2000/72849	WO	A1	2000-12-07	Hadasit Medical Research Services and Development		☐
	22	2003246780	JP	A	2003-09-02	Eisai	English Translation of Abstract from Japanese Patent Office	☐

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	1	ALFEREZ ET AL. "Inhibiting Signaling by erbB Receptor Tyrosine Kinases with AZD8931, a Potent Reversible small Molecule Inhibitor, Reduces Intestinal Adenoma Formation in the ApcMin/+ Mouse Model". EORTC-NCI-AACR (2010), Abstract 471	☐
	2	ALFEREZ ET AL. "Inhibiting Signaling by erbB Receptor Tyrosine Kinases with AZD8931, a Potent Reversible small Molecule Inhibitor, Reduces Intestinal Adenoma Formation in the ApcMin/+ Mouse Model". EORTC-NCI-AACR (2010), Poster	☐
	3	BLOWERS "AZD8931". IASLC ANNUAL TARGETED THERAPIES OF THE TREATMENT OF LUNG CANCER MEETING (2011), Santa Monica, CA, PowerPoint Presentation	☐
	4	CRISTOFANILLI ET AL. "Exploratory Subset Analysis According to Prior Endocrine Treatment of Two Randomized Phase II Trials Comparing Gefitinib (G) with Placebo (P) in Combination with Tamoxifen (T) or Anastrozole (A) in Hormone Receptor-Positive (HR+) Metastatic Breast Cancer (MBC)". J CLIN. ONCOL. (2009), Vol. 27, 15s, Abstract 1014	☐

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	5	HICKINSON ET AL. "AZD8931, an Equipotent, Reversible Inhibitor of Signaling by Epidermal Growth Factor Receptor, ERBB2 (HER2) and ERBB3: A Unique Agent for Simultaneous ERBB Receptor Blockade in Cancer". CLIN. CANCER RES. (2010), Vol. 16, No. 4, 1159-1169	☐
	6	KEILHOLZ ET AL. "Phase I Dose-Finding Study of Monotherapy with AZD8931, an Inhibitor of erbB1, 2 and 3 Signaling, in Patients with Advanced Solid Tumors". J CLIN ONCOL. (2011), Vol. 29, Abstract 3097	☐
	7	KEILHOLZ ET AL. "Phase I Dose-Finding Study of Monotherapy with AZD8931, an Inhibitor of erbB1, 2 and 3 Signaling, in Patients with Advanced Solid Tumors". ASCO (2011), Poster	☐
	8	KLINOWSKA ET AL. "AZD8931, an Equipotent, Reversible Inhibitor of erbB1, erbB2 and erbB3 Receptor Signaling: Characterisation of Pharmacological Profile". EUROPEAN JOURNAL OF CANCER SUPPLEMENTS (2009), Vol. 7, No. 2, 127	☐
	9	LOPEZ-MARTIN ET AL. "Phase I Dose-Finding Study of AZD8931, an Inhibitor of erbB1, 2 and 3 Receptor Signaling, in Combination with Paclitaxel". J CLIN. ONCOL. (2011), Vol. 29, Abstract 3105	☐
	10	LOPEZ-MARTIN ET AL. "Phase I Dose-Finding Study of AZD8931, an Inhibitor of erbB1, 2 and 3 Receptor Signaling, in Combination with Paclitaxel". ASCO (2011), Poster	☐
	11	MARSHALL ET AL. "Evaluation of AZD8931, an Equipotent Inhibitor of erbB1, erbB2 and erbB3 Receptor Signaling, on Ligand Stimulated Breast Cancer Cell Lines with Differing Levels of erbB2 Expression". SABCS (2009), Abstract 5059	☐
	12	NORMANNO ET AL. "Target-based therapies in breast cancer: current status and future perspectives". ENDOCR RELAT CANCER (2009), Vol. 16(3): 675-702.	☐
	13	SPEAKE ET AL. "Characterization of AZD8931, a Potent Reversible Small Molecule Inhibitor Against Epidermal Growth Factor Receptor (EGFR), Erythroblastic Leukemia Viral Oncogene Homolog 2 (HER2) and 3 (HER3) with a Unique and Balanced Pharmacological Profile". J CLIN. ONCOL. (2009), Vol. 27, 15s, Abstract 11072	☐

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